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Methane oxidation linked to chlorite dismutation

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We examined the potential for CH₄ oxidation to be coupled with oxygen derived from the dissimilatory reduction of perchlorate, chlorate, or via chlorite (ClO₂⁻) dismutation. Although dissimilatory reduction of ClO₄⁻ and ClO₃⁻ could be inferred from the accumulation of chloride ions either in spent media or in soil slurries prepared from exposed freshwater lake sediment, neither of these oxyanions evoked methane oxidation when added to either anaerobic mixed cultures or soil enriched in methanotrophs. In contrast, ClO₂⁻ amendment elicited such activity. Methane (0.2 kPa) was completely removed within several days from the headspace of cell suspensions of *Dechloromonas agitata* CKB incubated with either *Methylococcus capsulatus* Bath or *Methylobacterium album* BG8 in the presence of 5 mM ClO₂⁻. We also observed complete removal of 0.2 kPa CH₄ in bottles containing soil enriched in methanotrophs when co-incubated with *D. agitata* CKB and 10 mM ClO₂⁻. However, to be effective these experiments required physical separation of soil from *D. agitata* CKB to allow for the partitioning of O₂ liberated from chlorite dismutation into the shared headspace. Although a link between ClO₂⁻ and CH₄ consumption was established in soils and cultures, no upstream connection with either ClO₄⁻ or ClO₃⁻ was discerned. This result suggests that the release of O₂ during enzymatic perchlorate reduction was negligible, and that the oxygen produced was unavailable to the aerobic methanotrophs.

Keywords: chlorite, perchlorate, chlorate, methane, oxidation

INTRODUCTION

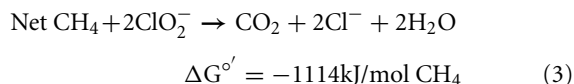
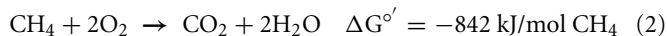
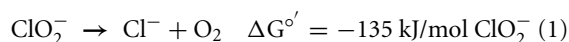
In-situ production of CO₂ by microbial activity is encouraged during enhanced oil recovery as a means of reducing oil viscosity and improving flow characteristics (Lazar et al., 2007; Youseff et al., 2009). Further, targeted growth of microbes and intentional precipitation of solid phase minerals can be applied to selectively decrease permeability and direct flow to enhance oil recovery (Jenneman et al., 1984; Zhu et al., 2013). These enhancements rely on the availability of appropriate electron acceptors to supply oxidant to microbes utilizing hydrocarbons or other reduced compounds as electron donors. Considerable attention has been paid thus far to the use of sulfate, nitrate or nitrite as electron acceptors in these applications (Youseff et al., 2009) with sulfate less favored because its reduction results in H₂S and leads to oil souring (Gieg et al., 2011). There has been recent interest in the use of perchlorate or chlorate, together known as (per)chlorate as electron acceptors. However, little is known about the fate of (per)chlorate in anoxic environments like oil reservoirs. Here we examine the potential reaction of (per)chlorate and chlorite with the low molecular weight hydrocarbon methane.

Chemical reduction of perchlorate is generally quite slow (Urbansky, 2002). However, under anoxic conditions dissimilatory perchlorate reducing bacteria (DPRB) rapidly reduce (per)chlorate to form chlorite. Chlorite thus formed is further degraded by these bacteria using chlorite dismutase to produce Cl⁻ and O₂ (Rikken et al., 1996; Kostan et al., 2010; Mlynek et al., 2011). These microbial processes reduce

(per)chlorate from both natural (Rao et al., 2007; Kounaves et al., 2010) and anthropogenic (Coates and Achenbach, 2004) sources. Degradation of intentionally added (per)chlorate is therefore likely in the proximity of hydrocarbon reservoirs given the abundance of suitable electron donors. There is a potential for O₂ liberation during this process (see reaction 1 below) that may be used by aerobic bacteria to oxidize aromatic compounds (benzene, naphthalene, catechol) via oxygenase-dependent pathways in otherwise anoxic soils and sediments (Coates et al., 1998, 1999a; Coates and Achenbach, 2004; Weelink et al., 2007; Carlström et al., 2013). A similar phenomenon was noted that could link biological oxidation of arsenite to the reduction of chlorate ions, presumably also by liberation of O₂ (Sun et al., 2010). This type of interaction has not been extended to the oxidation of low molecular weight hydrocarbons such as methane (CH₄).

Methane is produced by geothermal and microbial processes in the Earth's crust (Martini et al., 1996) and in marine and terrestrial sediments (Cicerone and Oremland, 1988). Methane is removed photochemically in the atmosphere by reaction with hydroxyl radicals but the most important removal mechanism in aqueous and terrestrial environments is by the action of anaerobic and aerobic methane oxidizing microbes (Cicerone and Oremland, 1988; Boetius et al., 2000). Significant quantities of methane are associated with oil reservoirs (Jones et al., 2007; Gieg et al., 2008) hence we hypothesize that ClO₂⁻ disproportionation, and by extension dissimilatory reduction of the upstream ClO₄⁻

or ClO_3^- ions, could be linked to aerobic CH_4 oxidation by a biochemical release of O_2 :



Two other well-studied microbiological processes can achieve a net oxidation of CH_4 under prevailing anaerobic conditions, (1) a reverse process of methanogenesis involving “ANME” archaea in syntrophy with bacterial sulfate- or sulfur-reduction (Hinrichs et al., 1999; Boetius et al., 2000; Milucka et al., 2012) and (2) nitrite-linked CH_4 oxidation that putatively liberates O_2 via NO dismutation as achieved by *Methyloirabilis oxyfera* (Ettwig et al., 2010). In our study we explored the potential for aerobic CH_4 oxidizing bacteria to utilize oxygen produced by DPRB during (per)chlorate reduction and chlorite dismutation.

MATERIALS AND METHODS

PREPARATION OF CULTURES

The DPRB *Dechloromonas agitata* CKB was grown at 30°C under N_2 on 20 mM sodium acetate and 10 mM NaClO_4 using phosphate buffer media (PBM) consisting of the following salts in solution (g/liter): Na_2HPO_4 (0.971), NaH_2PO_4 (0.379), NH_4Cl (0.25) plus 10 ml/l vitamins and 10 ml/l mineral stock solution (Sun et al., 2009). *Methylococcus capsulatus* Bath, *Methylosinus trichosporium* OB3b, and *Methyloirabilis albus* BG8, were grown and maintained at 30°C on air + 30 kPa CH_4 using nitrate mineral salts media (NMS; Whittenbury et al., 1970). Cultures (1 l) for washed cell suspensions were harvested during late exponential phase, centrifuged ($7000 \times g$), and washed twice with medium lacking substrates and vitamins. Final suspension volumes ranged from 5 to 150 ml. Cell concentrations at the start of incubations ranged from 1.8×10^8 cells ml^{-1} to 6.9×10^8 cells ml^{-1} .

PREPARATION OF SOILS AND SLURRIES

Soil from the seasonally exposed shoreline of Searsville Lake previously shown to harbor methanotrophic activity (Oremland and Culbertson, 1992) was air dried for two days at room temperature before sieving (<1 mm) to assure uniformity of soil particle size. Dried soil was stored for several weeks in stoppered 1 l glass flasks with air headspace and periodically augmented with 0.2 kPa CH_4 after consumption had removed all of the previously added CH_4 (4–6 days). Soil with thusly enhanced methanotrophic activity was used to determine CH_4 uptake in studies with added ClO_4^- and ClO_2^- in the absence of O_2 .

Sediment slurries were prepared by adding 100 ml SeFr2 freshwater media (flushed with 20 kPa CO_2 /80 kPa N_2 ; Miller et al., 2013) to 10 g Searsville Lake soil in N_2 flushed serum bottles (160 ml). Slurry pH was adjusted to 7.1 using 1 ml of 1 M NaHCO_3 . Slurries were incubated under N_2 headspace following periodic amendments with 1 to 2 mmoles acetate and 0.5 to 1 mmole ClO_4^- . Slurries were periodically sampled by syringe

using 22 g needles and filtered through a 0.2 μm Spin-X centrifuge tube. Acetate and ClO_4^- amendments were made after both were depleted (usually several days to weeks) during which time copious quantities of CH_4 were produced. Slurries with enhanced perchlorate reducing activity were used to determine methane uptake activity in the presence or absence of added O_2 .

MEASUREMENT OF (PER)CHLORATE REDUCTION AND CHLORITE DISMUTASE ACTIVITY

Aliquots of washed cell suspension of *D. agitata* CKB were distributed into stoppered and N_2 flushed 25 ml Balch tubes containing 10 ml PBM amended with 5 mM acetate and 10 mM NaClO_4 , NaClO_3 , or NaClO_2 . Initial cell densities were 1.8×10^8 cells per ml. The headspace was sampled over 7 days by syringe for CO_2 and O_2 . Aqueous samples (0.3 ml) were collected by syringe for analysis of dissolved acetate and anions (Cl^- , ClO_4^- , ClO_3^- , and ClO_2^-). A short-term (10 min) experiment was conducted to follow ClO_2^- disproportionation. In this study, triplicate samples were sacrificed at pre-determined times. Activity was stopped by addition of 0.1 ml 4N NaOH before measurement of headspace O_2 . Aqueous samples were subsequently collected for analysis of dissolved anions (Cl^- and ClO_2^-).

INCUBATIONS WITH MIXED CULTURES

Mixtures (10 ml) of *M. capsulatus* Bath and *D. agitata* CKB were prepared by adding washed cell suspensions of the cultures together in N_2 flushed Balch tubes (25 ml) sealed with butyl rubber stoppers. Methane (0.2 kPa) was introduced by syringe to all tubes and NaClO_2 (5 mM) was added to 3 tubes at the start of the incubation which was conducted at 37°C. Headspace CH_4 was monitored over 1 day. Single tubes were prepared without addition of ClO_2^- or without one of the cultures (i.e., no *D. agitata* CKB or no *M. capsulatus* Bath) to act as negative controls. A tube containing only *M. capsulatus* Bath under an air headspace acted as a positive control.

Additional microcosms were prepared in serum bottles (37 or 67 ml) using washed cell suspensions of *D. agitata* CKB and *M. trichosporium* OB3b or *M. albus* BG8. Inocula were either combined in bottles (5 ml each) in one aqueous phase or kept separate by placing methanotrophs (1 ml) inside an open-topped glass tube contained within the bottles before flushing with N_2 and later adding *D. agitata* CKB (5 ml) by syringe to the bottom of the bottles. In this manner, the cultures were segregated but shared a common headspace. Methane (0.2 kPa) was introduced by syringe to all bottles and NaClO_4 , NaClO_3 , or NaClO_2 (5 or 10 mM) was added aseptically to start the incubations which were conducted at 28°C. Headspace CH_4 and aqueous anions were monitored over time. Controls were prepared without additions of ClO_2^- .

INCUBATIONS WITH ^{14}C -LABELED CH_4

Washed cell suspensions (5 ml each) of *D. agitata* CKB and *M. trichosporium* OB3b were added together to N_2 flushed serum bottles (13 ml). Radiolabeled $^{14}\text{CH}_4$ (5 μCi ; specific activity = 21 $\mu\text{Ci}/\mu\text{mole}$) was added along with 1 kPa CH_4 to the headspace of each bottle. Perchlorate (5 mM) was added to triplicate bottles and ClO_2^- (5 mM) was added to a single bottle by syringe

to start the incubation which was conducted at 30°C. Gas samples for analysis of $^{14}\text{CH}_4$ and $^{14}\text{CO}_2$ were collected by syringe. At the end of the incubation, samples were acidified using 0.5 ml of 1.2N HCl to cause dissolved inorganic carbon ($\text{DIC} = \text{HCO}_3^- + \text{CO}_3^{2-}$) to react to form CO_2 gas which was partitioned into the headspace. The headspace was again sampled by syringe. Control incubations consisted of single bottles of *M. trichosporium* OB3b alone under an air headspace and *D. agitata* CKB alone under N_2 .

INCUBATIONS WITH SOIL SLURRIES

Slurry microcosms were prepared in N_2 flushed serum bottles (57 ml) containing 5 g of dried Searsville Lake soil with enhanced methanotrophic activity to which 10 ml Searsville Lake sediment slurry with enhanced perchlorate reducing activity (above) was added. Half the bottles were maintained under N_2 while half were flushed with air. Substrate (5 mM ClO_4^- or ClO_3^-) was added by syringe followed by 0.5 ml CH_4 (1 kPa). Incubations were conducted at 22°C. Headspace and liquid samples were collected by syringe over 9 days.

INCUBATIONS WITH CULTURES PLUS SOIL

Soil microcosms were prepared in serum bottles (67 ml) using washed cell suspensions of *D. agitata* CKB and dried Searsville Lake soil which was enhanced in methanotrophic activity (above). Soil (2 g) was placed inside open-topped glass tubes contained within the bottles prior to sealing and flushing with N_2 . Subsequently, *D. agitata* CKB (10 ml) was added by syringe to the bottom of the bottles followed by aseptic addition of 5 mM acetate. The culture and the soil were thus segregated under a common headspace. Methane (0.1 kPa) was introduced by syringe to all bottles and 10 mM NaClO_4 , NaClO_3 , or NaClO_2^- was added to *D. agitata* CKB to start the incubations. Incubations were conducted at 22°C. Headspace CH_4 and CO_2 and aqueous acetate and anions were monitored over 7 days.

ANALYTICAL

Headspace O_2 was determined by ECD-GC using a molecular sieve 5A column (3.2 mm O.D. \times 2.4 m) operated at 75°C using hydrocarbon-free UHP N_2 carrier. Background O_2 was minimized by flushing syringes and needles with O_2 -free N_2 prior to sampling. The detection limit was 0.05 mmol O_2/L . Headspace CH_4 and CO_2 were determined by FID- and TCD-GC, respectively (Miller et al., 2013). Cell densities were determined by direct cell counting of liquid samples using acridine orange epi-fluorescence microscopy (Hobbie et al., 1977). Additional aqueous samples, including slurries, were filtered using Spin-X centrifuge filter tubes (0.2 μm ; Corning Inc., Corning, NY) before determination of dissolved acetate by HPLC (Hoeft et al., 2004) or anions by IC (Miller et al., 2003). Dissolved ClO_4^- was analyzed separately by suppressed conductivity IC using a Dionex ISC 1100 containing an AS16 analytical column (4 \times 250 mm) and an AG16 guard column (4 \times 50 mm) with 0.035 M NaOH eluent. Measurements of headspace $^{14}\text{CH}_4$ and $^{14}\text{CO}_2$ were made by gas proportional counting (Culbertson et al., 1981) following TCD-GC analysis of CH_4 and CO_2 with separation on a Hayesep D column (100/120; 3.2 mm O.D. \times 4.8 m) using UHP He carrier.

CALCULATIONS

The total amount of gas in each bottle or tube was calculated from the headspace concentration using Henry's Law and the volumes of gas and liquid present. The dimensionless Henry's Law constants ($K_H = C_G/C_L$) used were 31.43 for O_2 , 29.46 for CH_4 and 1.20 for CO_2 and were not corrected for ionic strength.

RESULTS

(PER)CHLORATE REDUCTASE AND CHLORITE DISMUTASE ACTIVITY

Dissimilatory (per)chlorate reduction by *D. agitata* CKB resulted in conversion of 85–100 μmoles added ClO_4^- or ClO_3^- to Cl^- in the presence of 50 μmoles added acetate (Figures 1A,B). Much less ClO_4^- or ClO_3^- (<15 μmoles) was consumed without added acetate and a corresponding lesser amount of Cl^- was produced. These observations suggest endogenous metabolism of intrinsic electron donors such as glycogen or polyhydroxybutyrate (PHB). No activity was observed in killed controls or in incubations with media and chloroxyanions alone (data not shown). Biological reduction of ClO_4^- or ClO_3^- and consumption of acetate occurred over approximately 2 days. Chlorite dismutation was much more rapid. More than half of the 100 μmoles ClO_2^- added was consumed and converted to Cl^- before the initial sampling at $T = 2$ min (Figure 1C). An additional 20 μmoles ClO_2^- were consumed over 7 days, however more than 20 μmoles ClO_2^- remained unreacted at the end.

Carbon dioxide (CO_2) was the dominant gaseous product of dissimilatory reduction of ClO_4^- or ClO_3^- in the presence of acetate (Figures 2A,B). Slightly more CO_2 was produced than could be accounted for by the added acetate. Little CO_2 was produced without added acetate. Small amounts of O_2 (up to 15 μmoles) were produced during incubations with ClO_4^- or ClO_3^- . In contrast, ClO_2^- dismutation resulted in substantial and rapid O_2 production (Figure 2C) corresponding to release of >35% of the added ClO_2^- within the first day. As expected, there was no effect of added acetate on disproportionation of ClO_2^- ; however details of the early evolution of O_2 were obscured by the coarse sampling schedule.

The pattern of early O_2 production during ClO_2^- dismutation was made clear in a subsequent short-term (10 min) experiment where 56 μmoles of both O_2 and Cl^- were produced during the consumption of 56 μmoles of ClO_2^- (Figures 3A,B). A minor amount of CO_2 (<1 μmole) was produced (Figure 3C). Nearly half (40 μmoles) of the added ClO_2^- remained unreacted at the end of the experiment.

MIXED CULTURES OXIDIZED CH_4

Methane was oxidized by methanotrophic bacteria *M. capsulatus* Bath (Figure 4A) and *M. album* BG8 (Figure 4B) during the reaction of *D. agitata* CKB with ClO_2^- . No removal of CH_4 was observed in mixed cell suspensions amended with ClO_4^- or ClO_3^- (i.e., no added ClO_2^-). Methane consumption occurred while the methanotrophs were in direct contact with up to 10 mM ClO_2^- . Methane removal during incubations of co-cultures of *D. agitata* CKB with *M. capsulatus* Bath at 37°C occurred within 1 day while removal of CH_4 by *M. album* BG8 at 28°C occurred over 4 days. Similar rates of CH_4 consumption were observed for these mixed cultures whether they were segregated or co-mingled

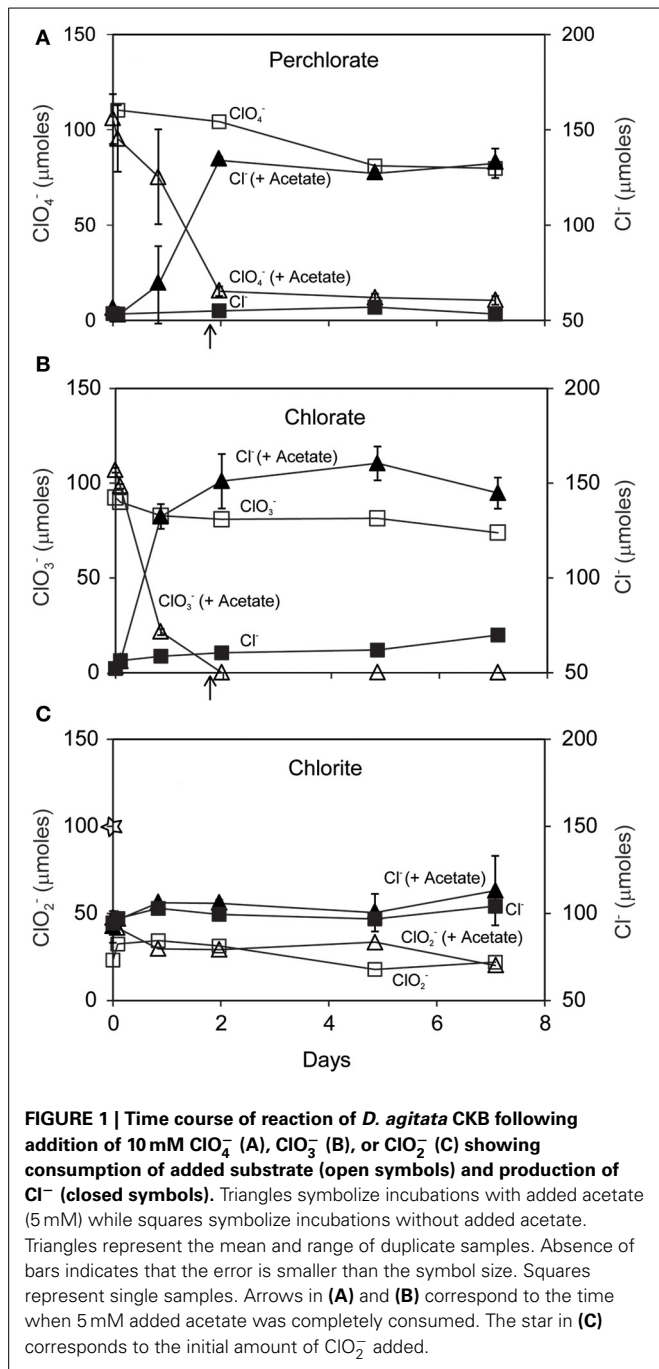


FIGURE 1 | Time course of reaction of *D. agitata* CKB following addition of 10 mM ClO_4^- (A), ClO_3^- (B), or ClO_2^- (C) showing consumption of added substrate (open symbols) and production of Cl^- (closed symbols). Triangles symbolize incubations with added acetate (5 mM) while squares symbolize incubations without added acetate. Triangles represent the mean and range of duplicate samples. Absence of bars indicates that the error is smaller than the symbol size. Squares represent single samples. Arrows in (A) and (B) correspond to the time when 5 mM added acetate was completely consumed. The star in (C) corresponds to the initial amount of ClO_2^- added.

(Figure 4B) indicating that direct contact of cells was not required for methane consumption to occur.

MIXED CULTURES OXIDIZE $^{14}\text{CH}_4$ TO $^{14}\text{CO}_2$

Strain *M. trichosporium* OB3b co-cultured with *D. agitata* CKB containing 5 mM acetate oxidized $^{14}\text{CH}_4$ directly to $^{14}\text{CO}_2$ (Figure 5). The rate of $^{14}\text{CH}_4$ loss under anaerobic conditions with 5 mM added ClO_2^- was similar to that under aerobic conditions. No loss of $^{14}\text{CH}_4$ occurred in mixed cell suspensions amended with 5 mM ClO_4^- in lieu of ClO_2^- or in controls without methanotrophs. During the incubation, the product of

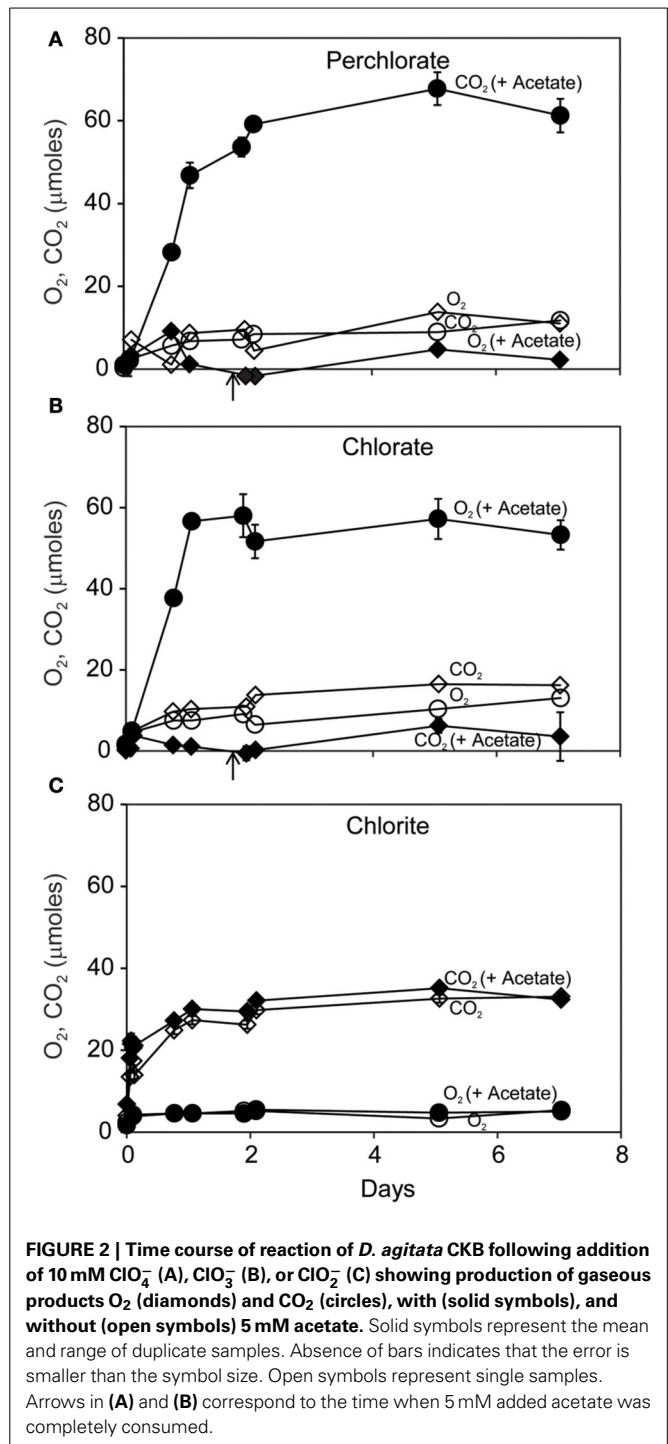
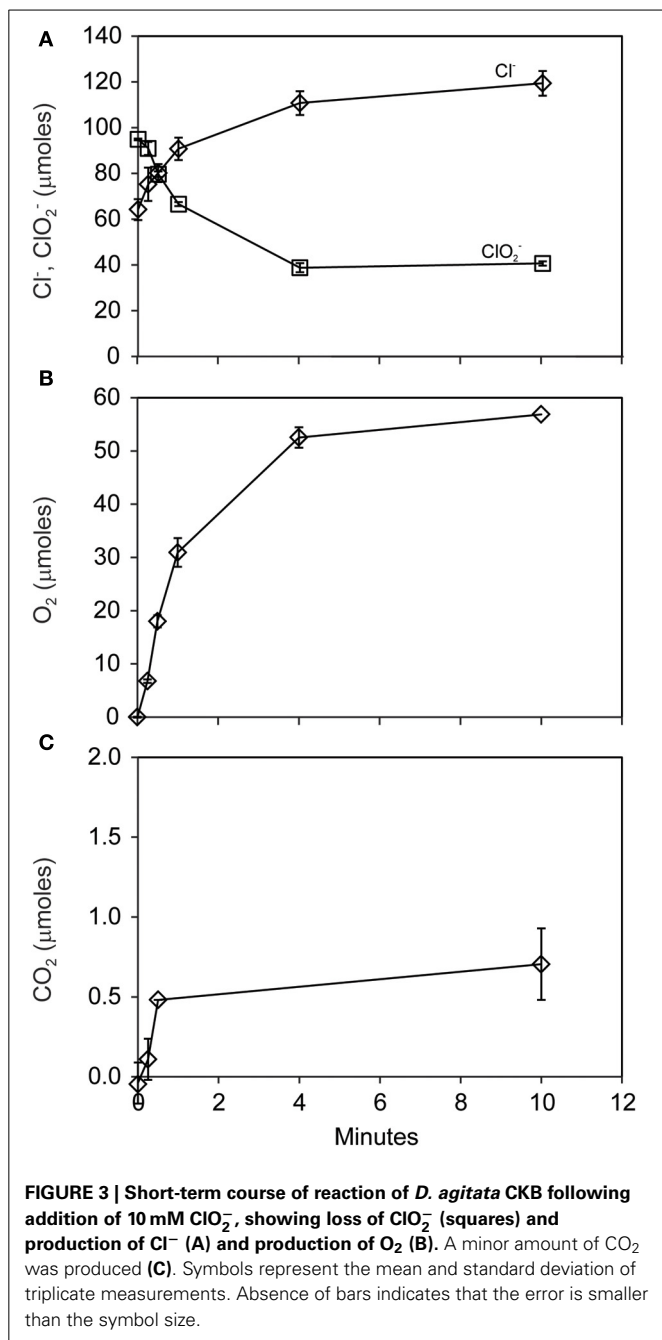


FIGURE 2 | Time course of reaction of *D. agitata* CKB following addition of 10 mM ClO_4^- (A), ClO_3^- (B), or ClO_2^- (C) showing production of gaseous products O_2 (diamonds) and CO_2 (circles), with (solid symbols) and without (open symbols) 5 mM acetate. Solid symbols represent the mean and range of duplicate samples. Absence of bars indicates that the error is smaller than the symbol size. Open symbols represent single samples. Arrows in (A) and (B) correspond to the time when 5 mM added acetate was completely consumed.

methanotrophy ($^{14}\text{CO}_2$) was distributed about equally between liquid and gas phases. Roughly 20% of the added $^{14}\text{CH}_4$ appeared as $^{14}\text{CO}_2$ in the headspace after 20 h (0.8 days). Most of the $^{14}\text{CH}_4$ added was recovered as $^{14}\text{CO}_2$ (60–90% recovery after acidification).

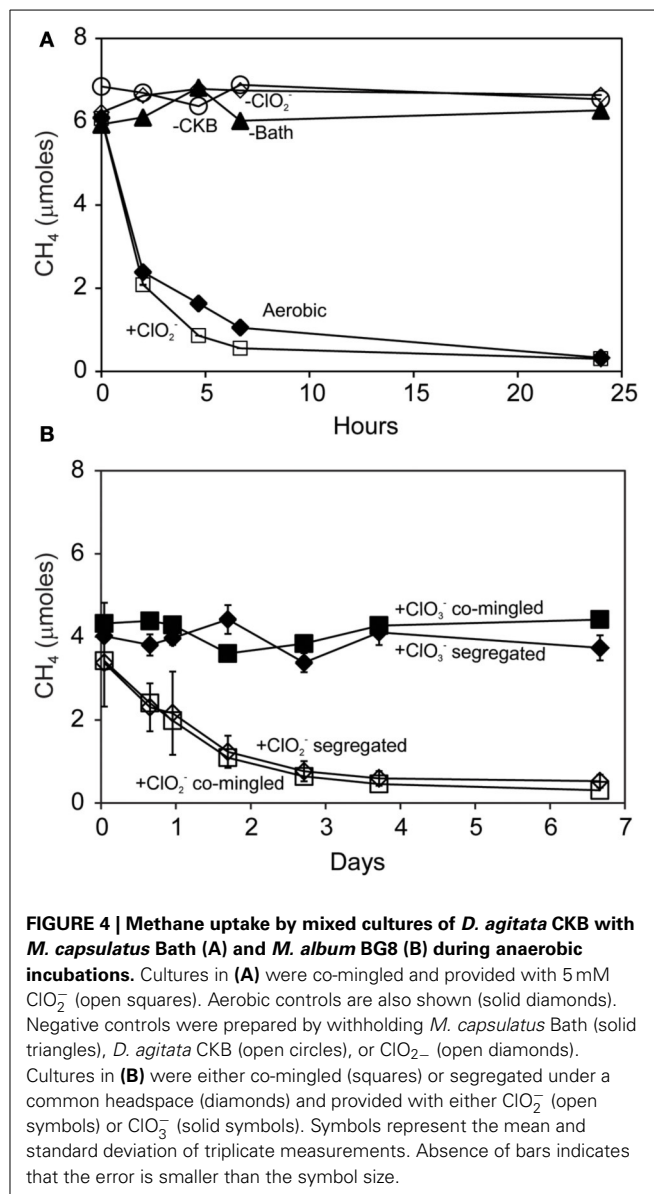
OXIDATION OF CH_4 BY SOILS

Searsville Lake sediment slurries removed repeated pulses of added ClO_4^- during anaerobic incubations using freshwater



media with added acetate. Over a period of 1 month, 4 additions of 10 mM ClO_4^- were removed in bottles with commensurate consumption of 4 additions of 5 mM added acetate (data not shown). These slurries, thus enhanced in ClO_4^- reducing capacity, were used in separate incubations with added CH_4 (Figure 6). Under aerobic conditions, incubations with or without additions of ClO_4^- or ClO_2^- (5 mM) completely consumed 30 μmoles CH_4 within 5 days. However, no oxidation of CH_4 occurred during anaerobic incubations of Searsville Lake sediment slurries either with or without additions of ClO_4^- or ClO_2^- .

Methane oxidation was observed when Searsville Lake soil (previously enhanced in methanotrophic activity) was segregated



from liquid cultures of *D. agitata* CKB. Methane was completely consumed over the next 5 days by soil methanotrophs during the reaction of DPRB with 10 mM ClO_2^- (Figure 7). No CH_4 loss was observed when DPRB were provided with either 10 mM ClO_4^- or ClO_3^- .

DISCUSSION

Few enzymes outside of photosystem II are capable of generating dioxygen in anoxic settings. In addition to chlorite dismutase (Cld), these include superoxide dismutase and catalase (McCord et al., 1971), and a putative nitric oxide dismutase (Ettwig et al., 2012). Chlorite dismutase has been purified from at least four DPRB (Mehboob et al., 2009) and is well studied (Coates et al., 1999b; Lee et al., 2008; Goblirsch et al., 2010, 2011). It is a heme enzyme which operates hyperspecifically; its assumed sole function is to detoxify ClO_2^- . The present study is the first report of anaerobic methane oxidation linked to the combined presence of DPRB and ClO_2^- . We emphasize that this is a “cryptic” aerobic methane oxidation in that the organisms oxidizing

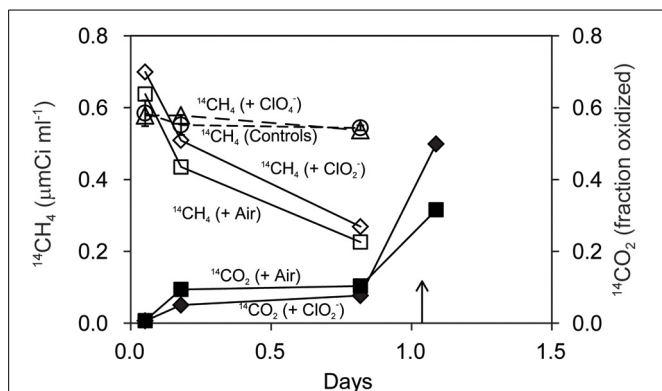


FIGURE 5 | Oxidation of $^{14}\text{CH}_4$ (open symbols) to $^{14}\text{CO}_2$ (closed symbols) by *M. trichosporium* OB3b during anaerobic incubations with *D. agitata* CKB following addition of ClO_4^- (triangles) and ClO_2^- (diamonds). Triangles represent the mean and standard deviation of triplicate measurements of samples with ClO_4^- added. All others were single bottles. Controls (no bacteria or no chloroxyanion added) were pooled and are shown as circles with error bars. Aerobic incubations with *M. trichosporium* OB3b alone are also shown (squares). Samples were acidified at the time indicated by the arrow.

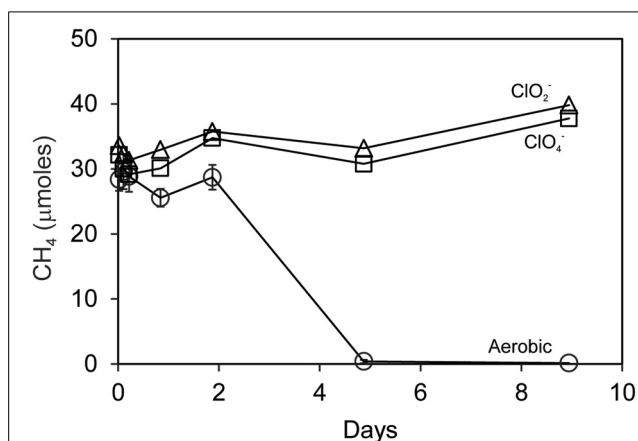


FIGURE 6 | Methane uptake by Searsville Lake sediment slurries pre-adapted to reduce perchlorate using acetate as electron donor. Anaerobic incubations with 5 mM added ClO_4^- (squares) or ClO_2^- (triangles) showed no uptake. Aerobic incubations (circles) represent the mean and standard deviation of CH_4 measurements in all bottles aerobic.

methane are aerobic methanotrophs as opposed to anaerobic methanogenic archaea utilizing reverse methanogenesis to consume methane (Hinrichs et al., 1999). As such, this process is somewhat analogous to nitrite-dependent anaerobic methane oxidation as purportedly carried out by the mixed culture containing *Methyloirabalis oxyfera* (Ettwig et al., 2010). In our study, methanotrophs use O_2 derived from disproportionation of ClO_2^- by DPRB (reaction 1) to oxidize CH_4 (reaction 3). This is in contrast to the mechanism proposed for *M. oxyfera* in which a single organism may use O_2 from disproportionation of NO derived from NO_2^- (reaction 4) to oxidize CH_4 (reaction 5):

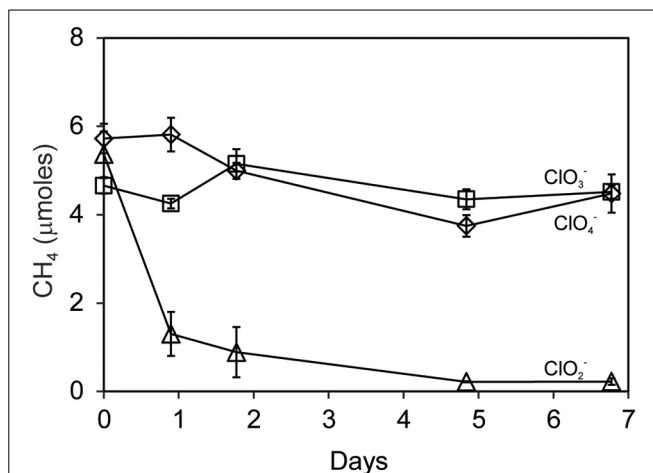
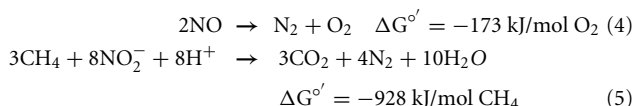


FIGURE 7 | Methane uptake by Searsville Lake soils during anaerobic incubations with *D. agitata* CKB following addition of ClO_2^- (triangles), ClO_4^- (diamonds), or ClO_3^- (squares). Symbols represent the mean and standard deviation of triplicate measurements. Absence of bars indicates that the error is smaller than the symbol size.

Previous work by Coates et al. (1998, 1999a); Coates and Achenbach (2006) demonstrated a link between Cld activity and other aerobic hydrocarbon oxidizing bacteria during the degradation of benzene and naphthalene. In their experiments with pristine soil and hydrocarbon contaminated sediment, ^{14}C -benzene was oxidized to $^{14}\text{CO}_2$ over several days when provided with ClO_2^- in the presence of washed cells of *D. agitata* CKB under anoxic conditions. Similarly, ^{14}C -naphthalene was rapidly oxidized to $^{14}\text{CO}_2$ in the presence of washed cells of *D. agitata* CKB and *Pseudomonas* sp. strain JS150 (an aerobic hydrocarbon oxidizer) when provided with ClO_2^- .

Here we demonstrated that O_2 released by the reaction of ClO_2^- with pure cultures of DPRB could be utilized by a variety of methane oxidizing bacteria, including γ -Proteobacteria (*M. capsulatus* Bath and *M. album* BG8) and α -Proteobacteria (*M. trichosporium* OB3b) methanotrophs. Addition of 10 mM ClO_2^- to DPRB resulted in only 40–60% recovery as O_2 and Cl^- (Figure 3). This may be attributed to a toxic effect of elevated ClO_2^- or bleaching of the Cld enzyme (Streit and DuBois, 2008). Nonetheless, much of the available O_2 was freely released during the reaction of ClO_2^- with *D. agitata* CKB, consistent with localization of Cld in the periplasm of DPRB (O'Connor and Coates, 2002). We demonstrated that direct addition of 5 or 10 mM ClO_2^- to mixed cultures of DPRB and methanotrophs did not inhibit the methanotrophs. We also showed that direct contact between the cells was not required as CH_4 oxidation also occurred when the cells were contained in separate compartments under a common headspace. We further showed that $^{14}\text{CH}_4$ was quantitatively oxidized to $^{14}\text{CO}_2$ by the methanotrophs in culture. Our conclusion is that methane oxidizers utilized O_2 provided by the dismutation of ClO_2^- by DPRB.

We were unable to link methane oxidation to perchlorate or chlorate reduction. Small amounts of oxygen were produced when cultures of *D. agitata* CKB were amended with ClO_4^- or

ClO_3^- (Figure 2), however methane was not consumed during co-culturing with methanotrophs (Figures 4, 6). This was previously observed for benzene and naphthalene (Coates et al., 1999b; Coates and Achenbach, 2006) and may be explained by O_2 scavenging attributable to other processes, including activation of a terminal oxidase during (per)chlorate reduction (Rikken et al., 1996). It is also possible that slower kinetics of ClO_4^- and ClO_3^- reduction limits the production and subsequent dismutation of ClO_2^- and therefore release of O_2 . Relief of this bottleneck could lead to more O_2 being available to aerobic methanotrophs and stimulation of the unique process described herein.

In-situ oxidation of CH_4 using O_2 derived from chlorite dismutation may be useful in removing elevated levels of CH_4 in subsurface environments. Chlorite is 10^4 times more soluble in water than O_2 and could be easily and safely directed to the anaerobic zone (where methane may be present) during bioremediation. One example is enhanced oxidation of landfill methane without the use of forced air, reducing the risk of fire and explosion. The ability of methanotrophs and DPRB to function in separate compartments under a common headspace could be exploited at distal stages of oil development, for instance at well heads where unusable CH_4 is typically flared off to reduce transportation costs or risk. In addition, production of CO_2 formed during oxidation of CH_4 may be viewed similarly to injected CO_2 in efforts to dissolve and flush oil from developed petroleum reservoirs (Blunt et al., 1993). Further, bioclogging by cells and biocementation resulting from carbonate precipitation may be enhanced by the growth and activity of microbes capable of linking methane oxidation with (per)chlorate reduction. Bioclogging and biocementation are features of microbial enhanced hydrocarbon recovery most likely to be exploited by geotechnologists to direct hydrocarbon flow into more permeable substrates in order to enhance recovery. However, it must be noted that we observed no methane oxidation in water saturated soils exposed to both CH_4 and ClO_2^- (Figure 6) indicating that O_2 derived from chlorite dismutation may face transport limitations in saturated anaerobic environments and may be consumed before reaching nearby CH_4 . Even under our unsaturated experimental conditions where liquid cultures and soil were segregated under a common headspace (Figure 7) the amount of soil present in each microcosm had to be optimized in order to balance methanotrophy with other soil O_2 utilizing processes.

Oxidation of CH_4 by the mechanism identified here may reduce the greenhouse impact of fugitive gases during hydrocarbon reservoir development and recovery because the global warming potential of CO_2 is 25 times lower than CH_4 on a 100 year time scale (IPCC 5th assessment, 2013). In addition, intermediates along the pathway of aerobic methane oxidation (e.g., methanol, formaldehyde, and formate) are themselves quite useful as chemical feedstocks for a myriad of industrial applications including biofuel production.

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